



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 767 179 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
27.06.2001 Bulletin 2001/26

(51) Int Cl.7: C08C 19/44, C08L 19/00,
C08K 3/36

(21) Application number: 96114951.5

(22) Date of filing: 18.09.1996

(54) Method for producing a rubber composition

Verfahren zur Herstellung einer Kautschukzusammensetzung

Procédé de préparation de composition de caoutchouc

(84) Designated Contracting States:
DE ES FR GB IT

(74) Representative:
Hansen, Bernd, Dr. Dipl.-Chem. et al
Hoffmann Eitle,
Patent- und Rechtsanwälte,
Arabellastrasse 4
81925 München (DE)

(30) Priority: 20.09.1995 JP 24190495

(56) References cited:
EP-A- 0 002 864 EP-A- 0 299 074
EP-A- 0 661 298

(43) Date of publication of application:
09.04.1997 Bulletin 1997/15

- PATENT ABSTRACTS OF JAPAN vol. 013, no. 479 (C-648), 18 October 1989 & JP 01 188501 A (BRIDGESTONE CORP; OTHERS: 01), 27 July 1989,

(73) Proprietor: Bridgestone Corporation
Tokyo 104-8340 (JP)

(72) Inventors:

- Toba, Tadashi, c/o Bridgestone Corp.
Tokyo 104 (JP)
- Shimalzu, Takashi, c/o Bridgestone Corp.
Tokyo 104 (JP)
- Kitamura, Takashi, c/o Bridgestone Corp.
Tokyo 104 (JP)

EP 0 767 179 B1

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

Description**BACKGROUND OF THE INVENTION****5 Field of the Invention**

[0001] The present invention relates to a method for producing a rubber composition into which a silica is incorporated, more specifically, to a method for producing a rubber composition which has a satisfactorily low hysteresis loss (low rolling resistance, low gas mileage) and well-balanced physical properties such as break characteristics, wear resistance and processability and which is suitable to use in a tire tread and the like.

Description of Related Art

[0002] Recently, tensile stress and break characteristics of silica-incorporating rubber compositions have been attempted to be improved. With development of increasingly advanced performance of tires, a rubber composition is required which has well-balanced physical properties such as processability and break characteristics as well as a satisfactorily low hysteresis loss for use as a tire tread rubber.

[0003] In an attempt to reduce the hysteresis loss, a modified rubber having a specific structure is employed, as a major component, in a carbon filler-containing rubber formulation. However, simultaneous achievement of high grip performance in a rainy day and low hysteresis loss is very difficult. In order to solve such problems, some rubber compositions have been known in which white carbon such as silica is employed as a filler in addition to carbon black, or only white carbon is employed.

[0004] For example, a rubber composition containing a polymer whose terminals are modified with a silane compound and a silica is reported (e.g., JP-B No. 52-5071, JP-A No. 56-104906, JP-A No. 62-50346, JP-A No. 62-227908). However, such polymer contains an insufficient amount of a residual alkoxy group and is a polymer in which only one side of the terminals is modified. Thus, the reinforcing effect and dispersing effect on fillers such as a silica, and eventually, the mechanical property improving effect are not satisfactory. Furthermore, such polymer, even when, for example, dilithium as an initiator is used (JP-A No. 62-227908), has both terminals modified with alkoxy silane. They have coupling capability, so that the molecular weight of the polymer becomes difficult to control practically and the viscosity of the polymer cement before and after modification becomes higher, resulting in problems with regard to steam strip resistance and time course change in the final bulk polymer.

[0005] On the other hand, there is also known a rubber composition containing a polymer whose terminals are modified with a nitrogen-containing compound, and a silica (e.g., JP-A No. 64-22940). According to this publication, in obtaining a diene polymer by using an alkyl lithium as an initiator, a monomer containing a nitrogen atom is used for copolymerization at the terminal side of polymerization, or the diene polymer is modified, at its terminal, with a compound such as aminobenzophenone. However, the resultant polymer is also a polymer in which only one side of the terminals is modified with the nitrogen-containing compound, similarly to those mentioned above. Thus, the reinforcing effect on the silica is unsatisfactory.

[0006] A rubber composition containing, together with a silica, a butadiene/styrene rubber which has a specific structure and which is obtained by solution polymerization is also provided (pneumatic tires, e.g., in JP-A No. 3-239737). This is not satisfactory, either, although the reinforcement by the silica is somewhat improved.

[0007] These polymers mainly contain diene polymers, which have low affinity with a silica and are difficult to be made into uniformly dispersed rubber compositions by means of kneading. To compensate this disadvantage, an expensive silane coupling agent is incorporated in a large amount, in most of the cases. Since the silane coupling agent is readily hydrolyzed in an atmosphere, it needs to be handled with care and is difficult to be made into rubber compositions with satisfactory reproducibility.

[0008] EP 0 002 864 A1 discloses a process for preparing polymers comprising the steps of polymerizing a monoalkenyl aromatic compound and/or a conjugated diene in the presence of a monoalkali metal compound to form an alkali metal terminated polymer and reacting the alkali metal terminated polymer with an alkoxy silane containing an epoxy group. Other ingredients, such as fillers, may be added to the polymer obtained by the above process.

[0009] EP 0 661 298 A2 describes a process for the manufacture of an elastomeric composition comprising the steps of copolymerizing a conjugated diene in the presence of n-butyllithium, modifying the obtained copolymer with an alkoxy silane having an epoxy group and mixing the functionalized copolymer with silica by milling.

[0010] Accordingly, the inventors previously concentrated their energies on developing the method for producing a rubber composition, and then the inventors previously proposed in JP-A No. 1-188501 (first published in 1979), a method for improving the properties of a silica-incorporating rubber composition in which a silica is mixed in a solvent with a polymer in which a specific modifier is employed. By using this method, affinity and dispersability of a diene polymer with the silica, break characteristics and wearing resistance are improved, as compared with the methods

using conventional silane coupling agents. Nevertheless, further improved properties of the rubber compositions for tire treads are demanded in response to increasingly advanced performance of recent automobiles.

SUMMARY OF THE INVENTION

5

[0011] The inventors have made much effort in modifying the above-mentioned method to provide a method for producing a rubber composition having various improved physical properties.

10

[0012] Accordingly, an object of the present invention is to provide a method for producing a rubber composition having a low hysteresis loss (low rolling resistance, low gas mileage) and having various physical properties such as wet grip property, break characteristics, wearing resistance and processability which are satisfactorily well-balanced.

15

[0013] The inventors have found that, by modifying a terminal of a specific polymer with a silane modifier having a specific functionality, a method can be obtained for producing a rubber composition having a low hysteresis loss (low rolling resistance, low gas mileage) and satisfactorily well-balanced physical properties. As a result the present invention has been established.

15

[0014] Thus, a method for producing a rubber composition according to the present invention comprises:

the step wherein a conjugated diene monomer, or a conjugated diene monomer/a vinyl aromatic hydrocarbon monomer are polymerized by using an organic lithium initiator;

the step wherein an active terminal of the polymer or copolymer thus obtained is modified with a modifier represented by the general formula (I) shown below and having reactivity with a silica surface; and,

the step wherein the terminally-modified polymer or terminally-modified copolymer thus obtained is mixed with a silica in an organic solvent.

25



wherein R^1 denotes a substituent having an epoxy group or an unsaturated carbonyl group, X denotes an alkyl group or a halogen atom, R^2 denotes a group selected from the group consisting of aliphatic, alicyclic and aromatic hydrocarbons of 1 to 20 carbon atoms, m denotes an integer from 1 to 3, n denotes an integer from 1 to 3 and $m+n$ is an integer from 2 to 4.

30

[0015] The achievement of the above-mentioned object is possibly due to reactivity of the filler silica surface with the alkoxy functionality supplied by the modifier to the polymer, as well as due to high affinity of hydroxyl groups formed by the reaction of the substituted epoxy group or unsaturated carbonyl group with silanol groups on the silica surface.

35

PREFERRED EMBODIMENTS OF THE INVENTION

40

[0016] In the first step of the method according to the present invention, a conjugated diene monomer or a conjugated diene monomer/a vinyl aromatic hydrocarbon monomer are polymerized using an organic lithium initiator. The conjugated diene monomer may be a conjugated diene hydrocarbon having in its molecule 4 to 12 carbon atoms, preferably 4 to 8 carbon atoms. For example, 1,3-butadiene, isoprene, 2,3-dimethyl-1,3-butadiene, 1,3-pentadiene, octadiene and the like may be employed alone or in a mixture thereof, and 1,3-butadiene and isoprene are particularly preferred.

[0017] The vinyl aromatic hydrocarbon monomer for forming a copolymer together with the conjugated diene monomer includes, for example, styrene, α -methylstyrene, p -methylstyrene, o -methylstyrene, p -butylstyrene, vinyl naphthalene and the like, and styrene is particularly preferred.

45

[0018] The organic lithium compound used herein as an initiator may be alkyl lithium such as n -butyl lithium, sec -butyl lithium, ethyl lithium, propyl lithium, t -butyl lithium and hexyl lithium, alkylene dilithium such as 1,4-dilithiobutane, other lithiohydrocarbon such as phenyl lithium, stilbene dilithium and reaction products of butyl lithium and divinyl benzene, or organic lithiometals such as lithium tributyltin, lithium amides such as lithium diethylamide, N -methylbenzyl lithium amide, diethyltin amide, lithium piperidide, lithium pyrrolide and lithium hexamethylene imide, and organic lithium compounds including tertiary amine lithiums such as dimethylaminopropyl lithium and diethylaminopropyl lithium.

50

[0019] These organic lithium initiators may be employed alone or in a mixture thereof. The method for initiating polymerization using the organic lithium initiator may be a method wherein the polymerization is initiated by adding a monomer to a system containing a lithium amide initiator obtained from a secondary amine compound such as diethylamine or an imine compound such as hexamethylene imine and an organic lithium compound, a method wherein the polymerization is proceeded by adding a secondary amine compound or an imine compound to a system containing a monomer and an organic lithium compound to obtain a lithium amide catalyst as an initiator, or a method wherein the polymerization is initiated using only an organic lithium compound (for example butyllithium) without using a sec-

ondary amine compound or an imine compound.

[0020] The secondary amine compound used together with the lithium initiator includes, for example, dimethylamine, diethylamine, dipropylamine, di-n-butylamine, diisobutylamine, dipentylamine, dihexylamine, diheptylamine, dioctylamine, diallylamine, dicyclohexylamine, butylisopropylamine, dibenzylamine, methylbenzylamine, methylhexylamine and ethylhexylamine. Among them, an amine having a group selected from aliphatic hydrocarbon groups of 1 to 10 carbon atoms is preferable in view of further improvement in break characteristics and low hysteresis loss.

[0021] The imine compound mentioned above includes, for example, aziridine, azetidine, pyrrolidine, piperidine, 2-methylpiperidine, 3-methylpiperidine, 4-methylpiperidine, 3,5-dimethylpiperidine, 2-ethylpiperidine, hexamethylene-imine, heptamethyleneimine, coniine, morpholine, N-methylpiperazine, N-ethylpiperazine, N-methylimidazoline, N-ethylimidazoline, oxazine, pyrrolidine, pyrrole and azepine. Among them, a group selected from imine compound residue of 4 to 12 carbon atoms is preferable in view of further improvement in break characteristics and low hysteresis loss.

[0022] Among the initiators listed above, n-butyllithium or sec-butyllithium is a preferred alkylolithium in view of solubility in hydrocarbon solvents and initiation rate, for example, in cases where such alkylolithium is employed as an initiator, simultaneously with a secondary amine compound or imine compound. On the other hand, in cases where a lithiumamide initiator is employed from the beginning, lithium hexamethyleneimide, lithium pyrrolidine and lithium diethylamide are preferred in view of solubility in hydrocarbon solvents and reinforcing effect on a silica.

[0023] The amount of the lithium compound is 0.2 to 30 mmol per 100 g of a monomer. An amount less than 0.2 mmol or more than 30 mmol results in a polymer having unsatisfactory processability.

[0024] A conjugated diene polymer obtained in this step is preferably polybutadiene, which has an excellent wearing resistance. A conjugated diene/vinyl aromatic hydrocarbon copolymer is preferably butadiene/styrene copolymer, which has an excellent wearing resistance and aging resistance.

[0025] The microstructure (i.e., cis-1,4-, trans-1,4, vinyl) of the butadiene moiety of a conjugated diene copolymer, for example, polybutadiene and butadiene/styrene copolymer is not particularly limited; However, it may usually be one formed when an organic lithium catalyst system is used.

[0026] The concentration of a monomer in a solvent is usually 5 to 50 % by weight, preferably 10 to 35 % by weight. In a case of a copolymer of a conjugated diene and a vinyl aromatic hydrocarbon, the vinyl aromatic hydrocarbon content in the monomer mixture charged is 5 to 70 % by weight, preferably 10 to 50 % by weight.

[0027] Although the composition or ratio of a conjugated diene/vinyl aromatic hydrocarbon copolymer, for example, butadiene/styrene copolymer, is not particularly limited, a weight ratio of 30/70 to 95/5 is usually employed. The composition distribution of the copolymer may be a random distribution, a block distribution or an intermediate distribution between them, with a random or intermediate distribution being usually preferred.

[0028] In the second step in the method for producing according to the present invention, the polymer or copolymer (hereinafter referred to as (co)polymer, as appropriate) obtained as described above is modified with a modifier represented by the general formula (I) shown below.

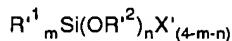


[0029] The modifier represented by the general formula (I) shown above is a silane compound having an epoxy or glycidoxy group or equivalent, and in the formula R^1 is a substituent having an epoxy group or an unsaturated carbonyl group. Preferably the number of carbon atoms of R^1 is from 3 to 20.

[0030] X denotes an alkyl group (preferably of 1 to 20 carbon atoms) or halogen atom, and R^2 denotes a group selected from aliphatic, alicyclic and aromatic hydrocarbon groups of 1 to 20 carbon atoms. R^2 is preferably a group selected from aliphatic, alicyclic and aromatic hydrocarbon groups of 1 to 13 carbon atoms. The aliphatic and alicyclic hydrocarbon groups are preferably those selected from aliphatic and alicyclic hydrocarbon groups, each of which has 1 to 6 carbon atoms in view of further improvement in break characteristics and low hysteresis loss. An alkyl group of 1 to 3 carbon atoms is more preferred in view of reactivity with a silica surface. The aromatic hydrocarbon group may be, for example, phenyl, naphthyl, biphenyl, anthryl and phenanthryl, with phenyl being preferred. Among those described above, the most preferable group is that selected from methyl, ethyl, propyl and phenyl, which are capable of further improving break characteristics and low hysteresis loss.

[0031] m denotes an integer from 1 to 3, n denotes an integer from 1 to 3 and $m+n$ is an integer from 2 to 4. m and n may vary within the range meeting the above-mentioned provision depending on a production condition and are not particularly limited. Both of a pure compound whose "m" and "n" are fixed and a mixture of compounds whose "m"s and "n"s are different to each other may be used in the method according to the present invention.

[0032] A preferred formula of the general formula (I) is the following general formula (Ia):



wherein R¹ denotes a group selected from the group consisting of γ -glycidoxethyl, γ -glycidoxypropyl, γ -glycidoxbutyl, γ -methacryloxyethyl, γ -methacryloxyethyl, γ -methacryloxypropyl, β -(3,4-epoxycyclohexyl)ethyl and β -(3,4-epoxycyclohexyl)propyl, X' denotes an alkyl group of 1 to 20 carbon atoms or a halogen atom, R² denotes a group selected from the group consisting of methyl, ethyl, propyl, butyl, phenyl, naphthyl, biphenyl, anthryl and phenanthryl, m denotes an integer from 1 to 3, n denotes an integer from 1 to 3 and m+n is an integer from 2 to 4.

[0033] The modifier represented by the general formula (I) includes, for example, γ -glycidoxethyl trimethoxsilane, γ -glycidoxypropyl trimethoxsilane, γ -glycidoxbutyl trimethoxsilane, γ -glycidoxypropyl triethoxysilane, γ -glycidoxypropyl tripropoxysilane, γ -glycidoxypropyl tributoxysilane, γ -glycidoxypropyl triphenoxysilane, γ -glycidoxypropyl methyldimethoxsilane, γ -glycidoxypropyl ethyldimethoxsilane, γ -glycidoxypropyl ethyldiethoxysilane, γ -glycidoxypropyl methyldiethoxysilane, γ -glycidoxypropyl methylpropoxysilane, γ -glycidoxypropyl methyl dibutoxysilane, γ -glycidoxypropyl methyl diphenoxysilane, γ -glycidoxypropyl dimethylmethoxysilane, γ -glycidoxypropyl diethylethoxysilane, γ -glycidoxypropyl dimethylethoxysilane, γ -glycidoxypropyl dimethylphenoxy silane, γ -glycidoxypropyl diethylmethoxysilane, γ -glycidoxypropyl methyl diisopropanoxysilane, bis(γ -glycidoxypropyl)dimethoxsilane, bis(γ -glycidoxypropyl)diethoxysilane, bis(γ -glycidoxypropyl)dipropoxysilane, bis(γ -glycidoxypropyl)dibutoxysilane, bis(γ -glycidoxypropyl)diphenoxysilane, bis(γ -glycidoxypropyl)methylmethoxysilane, bis(γ -glycidoxypropyl)methylethoxysilane, bis(γ -glycidoxypropyl)methylpropoxysilane, bis(γ -glycidoxypropyl)methylbutoxysilane, bis(γ -glycidoxypropyl)methylphenoxy silane tris(γ -glycidoxypropyl)methoxysilane, γ -methacryloxypropyl trimethoxysilane, γ -methacryloxypropyl triethoxysilane, γ -methacryloxyethyl trimethoxysilane, γ -methacryloxyethyl triethoxysilane, bis(γ -methacryloxypropyl)dimethoxysilane, tris(γ -methacryloxypropyl)methoxysilane, β -(3,4-epoxycyclohexyl)ethyl-trimethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-triethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-tripropoxysilane, β -(3,4-epoxycyclohexyl)ethyl-tributoxysilane, β -(3,4-epoxycyclohexyl)ethyl-triphenoxysilane, β -(3,4-epoxycyclohexyl)propyl-trimethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldimethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-ethyldimethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-ethyl diethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldiethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldipropoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldibutoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldiphenoxysilane, β -(3,4-epoxycyclohexyl)ethyl-dimethylmethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-diethylethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-dimethylethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-dimethylpropoxysilane, β -(3,4-epoxycyclohexyl)ethyl-dimethylbutoxysilane, β -(3,4-epoxycyclohexyl)ethyl-dimethylphenoxy silane, β -(3,4-epoxycyclohexyl)ethyl-diethylmethoxysilane, β -(3,4-epoxycyclohexyl)ethyl-methyldiisopropanoxysilane and the like. Among them, γ -glycidoxypropyl trimethoxysilane and γ -methacryloxypropyl trimethoxysilane are preferable since they further improve the physical properties such as wearing resistance as well as low hysteresis loss.

[0034] The amount of a modifier used is 0.1 to 10 molecules, preferably 0.2 to 3 molecules per one lithium atom. An amount departing from the range specified above results in reduced reaction efficiency of the silica, which may cause poor dispersion of the silica.

[0035] The polymerization reaction and the reaction between the polymeric active terminals and a modifier, described above, may be conducted at a constant temperature or temperature being elevated gradually, preferably within the range from 0 to 150 °C.

[0036] The polymerization reaction is conducted by allowing a monomer in a liquid phase to be in contact with a catalyst under a pressure sufficient to keep the liquid phase. All materials to be included in the reaction system are preferably free from the substances interfering the catalytic effect. The reaction conducted with an added modifier after polymerization should be maintained in an atmosphere in which the active terminals of a (co)polymer are not in activated.

[0037] In this method, the organic solvent employed as a polymerization solvent includes aromatic hydrocarbon solvents such as benzene, toluene and xylene, aliphatic hydrocarbon solvents such as n-pentane, n-hexane, n-butane, 1-methylbutene, heptane and octane, alicyclic hydrocarbon solvents such as methylcyclopentane and cyclohexane and mixture thereof. Any of solvents capable of dissolving or dispersing the (co)polymer mentioned above uniformly may be employed without particular limitation.

[0038] In this polymerization, when a (co)polymer having a specific molecular structure is desired, a randomizer is preferably employed. The randomizer herein means a compound having the functions to control the microstructure of the conjugated diene polymer, for example, to increase the rate of 1,2-bonds in the butadiene moiety of a butadiene polymer or butadiene/styrene copolymer, or the ratio of 3,4-bonds of an isoprene polymer and to control the composition distribution of the monomer units of a conjugated diene/a vinyl aromatic hydrocarbon copolymer, for example, to randomize the butadiene units and styrene units of a butadiene/styrene copolymer.

[0039] The randomizer which may be used in the present invention is not particularly limited, and all those employed usually may be included. Examples of the randomizers are listed below.

- 55
 - (1) Ethers
 - (2) o-Dimethoxybenzenes
 - (3) Complexes of an alkaline metal with a ketone or triphosphite

(4) Compounds represented by the general formula shown below



5

wherein R is selected from aliphatic, alicyclic and aromatic hydrocarbons, M¹ is an alkaline metal, particularly lithium, sodium, potassium, rubidium or cesium, M² is an alkaline earth metal, typically calcium or barium, and n is an integer of 1 to 3.

(5) Tertiary amines

10

[0040] The randomizers are detailed below and they may be employed alone or in a mixture thereof.

(1) Examples of the ethers include 1,2-dimethoxyethane, 1,2-diethoxyethane, tetrahydrofuran, 2-methoxyethyl-tetrahydrofuran, diethylether, triethyleneglycol dimethyl ether and the like.

(2) Examples of the o-dimethoxybenzenes are veratrol and isohomoveratrol.

(3) Examples of a complex of an alkaline metal with a ketone or triphosphite are complexes of lithium, sodium, potassium, rubidium and cesium with a ketone such as acetone, methyl ethyl ketone, disopropyl ketone, benzophenone, acetophenone, dibenzyl ketone, fluorenone, xanthone, Michler's ketone and acetylacetone or with a triphosphite such as triethyl phosphite, trioctyl phosphite, tribenzyl phosphite and trinonyl phosphite.

(4) The randomizers represented by the general formula are detailed below.

Examples of alkaline metal salts or alkaline earth metal salts of alcohols and phenols, represented by the general formulas R(OM¹)_n and (RO)₂M² are lithium, sodium, potassium, rubidium, cesium, calcium and barium salts of methyl alcohol, ethyl alcohol, isopropyl alcohol, t-butyl alcohol, t-amyl alcohol, cyclohexyl alcohol, allyl alcohol, 2-butetyl alcohol, benzyl alcohol, phenol, catechol, resorcinol, hydroquinone, 1-naphthyl alcohol, p-nonyl phenol, pyrogallol and the like.

Examples of the carboxylate and acidic carbonate salts of the alkaline metals, represented by the general formulas R(COOM¹)_n and ROCOOM¹ are lithium, sodium, potassium, rubidium and cesium salts of isovaleric acid, lauric acid, palmitic acid, stearic acid, oleic acid, rosin acid, benzoic acid, pimelic acid, acidic n-dodecyl carbonate and acidic phenyl carbonate.

Examples of the sulfonates and sulfates of the alkaline metals, represented by the general formulas RSO₃M¹ and ROSO₃M¹ include lithium, sodium, potassium, rubidium and cesium salts of dodecylbenzene sulfonate, diisopropylnaphtharene sulfonate, N-methyl-N-methanesulfonate laurylamine, sulfate salt of lauryl alcohol and caproyl ethylene glycol sulfate.

(5) Examples of tertiary amines are triethylamine and tetramethyl ethylenediamine.

35

[0041] Among them, preferable randomizers are ethers in section (1) and R(OM¹)_n compounds in section (4) listed above, especially with which the molecular structure of a rubber composition obtained by the present method can readily be controlled.

[0042] The amount of a randomizer used is within the range from 0.01 to 1000 molar equivalents per molar equivalent of an organic lithium compound.

[0043] In the third step in the method according to the present invention, after completion of modification reaction of the terminally-modified (co)polymer obtained mentioned above, the (co)polymer allowed to remain in the above-mentioned organic solvent, i.e., the polymerization solvent, or the polymerization solvent is removed and the organic solvent listed above is newly added and the terminally-modified (co)polymer is dissolved therein. Into the system, a silica in a required amount is added. In the method according to the present invention, it is important to mix the silica with the (co)polymer in the presence of the organic solvent, in view of uniform dispersion of the silica and reactivity with the (co)polymer.

[0044] The silica employed in the method for producing a rubber composition according to the present invention includes, for example, silicon compounds such as wet silica (hydrated silicic acid), dry silica (anhydrous silicic acid), calcium silicate and aluminium silicate. A wet silica is preferable since it provides improvement in break characteristics while providing both of high grip property and low rolling resistance simultaneously.

[0045] The amount of a silica incorporated is 10 to 100 parts by weight per 100 parts by weight of the terminally-modified (co)polymer described above, and preferably 20 to 60 parts by weight in view of the reinforcement by the silica and efficiency of accompanying improvement in various physical properties. An amount less than 10 parts by weight results in unsatisfactory break characteristics, while an amount exceeding 100 parts by weight causes poor processability.

[0046] Mixture of the silica with the terminally-modified (co)polymer is conducted by reacting the (co)polymer with the silica well, at a temperature of 0 to 150 °C, preferably 50 to 130 °C.

[0047] A filler to be added to a rubber composition produced by the method according to the present invention may exclusively be the silica incorporated herein. In such case, the silica is employed in an amount of 10 to 100 parts by weight based on 100 parts by weight of the entire rubber material containing the terminally-modified (co)polymer, and preferably 20 to 60 parts by weight, in view of reinforcement and accompanying improvement in various physical properties. An amount less than 10 parts by weight results in unsatisfactory break characteristics, while an amount exceeding 100 parts by weight causes poor processability.

[0048] Alternatively, a carbon black may also be employed as a filler in addition to the silica. In such case, enhanced improvement in various physical properties can be obtained. The carbon black to be employed includes carbon blacks such as FEF, SRF, HAF, ISAF, SAF and the like, and preferably those having a iodine absorption (IA) of 60 mg/g or more and dibutylphthalate oil absorption value (DBP) of 80 ml/100 g or more. HAF, ISAF and SAF having an excellent wearing resistance are particularly preferred. The amount of a carbon black to be incorporated is not particularly limited provided that the effect of the silica is exhibited. It is however preferable in view of reinforcement and processability to incorporate from 0.1 to 90 parts by weight of the carbon black and from 10 to 99.9 parts by weight of the silica into 100 parts by weight of a starting rubber.

[0049] When mixing the silica with the above-mentioned (co) polymer, a filler such as the carbon black and a process oil may be added to the mixture in the presence of organic solvents.

[0050] The process oil which may be employed in the present invention includes, for example, paraffin oils, naphthene oils and aromatic oils. The aromatic oils are used in cases where high tensile strength and wearing resistance are important, while the naphthene and paraffin oils are used in cases where low hysteresis loss and low temperature characteristics are important. The amount used is 0 to 100 parts by weight per 100 parts by weight of the (co)polymer. An amount exceeding 100 parts by weight markedly affects the tensile strength and low hysteresis loss of a vulcanized rubber finally obtained adversely.

[0051] After mixing and reacting with the silica, the solvent is removed to obtain a master batch rubber composition according to the present invention. The solvent may be removed, for example, by purging with steam, by adding a poor solvent such as methanol into the reaction system to solidify the polymer followed by drying with a thermal roll or under reduced pressure, or by subjecting the polymer solution directly to the thermal roll or reduced pressure, according to conventional procedures.

[0052] The Mooney viscosity (ML₁₊₄, 100 °C) of the master batch rubber composition thus obtained is preferably within the range from 20 to 200. A Mooney viscosity less than 20 results in poor tensile strength and affects break characteristics adversely, while that exceeding 200 causes problems with regard to processability.

[0053] The master batch rubber composition thus obtained according to the present method may be optionally blended with natural rubbers or other synthetic rubbers, together with other additives and vulcanizing agents, as desired.

[0054] When blending with the natural rubbers or other synthetic rubbers, the master batch rubber composition should be contained in an amount of 10 parts by weight or more per 100 parts by weight of the entire rubber material employed for the rubber composition, and preferably in an amount of 40 parts by weight or more for the purpose of realizing the advantageous effect of the polymer according to the present invention sufficiently. For example, in a blend with a natural rubber, the polymer obtained by the present invention in an amount less than 10 parts by weight affects the balance of necessary properties of a rubber composition adversely.

[0055] The synthetic rubber which may be blended is cis-1,4-polyisoprene, styrene/butadiene copolymer, low cis-1,4-polybutadiene, high cis-1,4-polybutadiene, ethylene/propylene/diene copolymer, chloroprene, halogenated butyl rubber, acrylonitrile/butadiene rubber (NBR) and the like. Among them, it is preferable to blend a natural rubber, high cis-1,4-polybutadiene and halogenated butyl rubber in view of tensile strength, wearing resistance and processability.

[0056] In the rubber composition obtained according to the present method, a silane coupling agent is incorporated in order to further improve the reinforcing effect of a silica. The examples of the silane coupling agent are listed below.

Bis(3-triethoxysilylpropyl)tetrasulfide, bis(2-triethoxysilylethyl)tetrasulfide, bis(3-trimethoxysilylpropyl)tetrasulfide, bis(2-trimethoxysilylethyl)tetrasulfide, 3-mercaptopropyltrimethoxysilane, 3-mercaptopropyltriethoxysilane, 2-mercaptopoethyltrimethoxysilane, 2-mercaptopoethyltriethoxysilane, 3-nitropropyltrimethoxysilane, 3-nitropropyltriethoxysilane, 3-chloropropyltrimethoxysilane, 3-chloropropyltriethoxysilane, 2-chloroethyltrimethoxysilane, 2-chloroethyltriethoxysilane, 3-trimethoxysilylpropyl-N,N-dimethylthiocarbamoyltetrasulfide, 3-triethoxysilylpropyl-N,N-dimethylthiocarbamoyltetrasulfide, 2-triethoxysilylethyl-N,N-dimethylthiocarbamoyltetrasulfide, 3-trimethoxysilylpropylbenzothiazole tetrasulfide, 3-triethoxysilylpropylbenzothiazole tetrasulfide, 3-triethoxysilylpropylmethacrylate monosulfide, 3-trimethoxysilylpropylmethacrylate monosulfide and the like.

[0057] Bis(3-triethoxysilylpropyl)tetrasulfide and 3-trimethoxysilylpropylbenzothiazole tetrasulfide are preferred in view of the reinforcement improving effect. Other examples are bis(3-dieethoxymethylsilylpropyl)tetrasulfide, 3-mercaptopropyltrimethoxymethylsilane, 3-nitropropyltrimethoxysilane, 3-chloropropyltrimethoxymethylsilane, dimethoxymethylsilylpropyl-N,N-dimethylthiocarbamoyltetrasulfide and dimethoxymethylsilylpropylbenzothiazole tetrasulfide.

[0058] When a silane coupling agent is incorporated in a rubber composition obtained according to the present method, the amount of the agent may be smaller than a conventional amount for maintaining the equivalent physical

properties. In view of the reinforcement, the amount may be 0.1 to 10 parts by weight, preferably 0.1 to 5 parts by weight per 100 parts by weight of the rubber material although it may vary depending on the amount of the silica incorporated

[0059] A vulcanizing agent employed in a rubber composition obtained according to the present invention may be 5 sulfur and the like, which is employed in an amount of 0.1 to 5 parts by weight, preferably 1 to 2 parts by weight per 100 parts by weight of the rubber material. An amount less than 0.1 parts by weight results in affecting break strength, wearing resistance and hysteresis loss of a vulcanized rubber adversely, while an amount exceeding 5 parts by weight causes loss of rubber elasticity.

[0060] A vulcanization accelerator which can be employed in the present invention is not particularly specified, and 10 may preferably be thiazoles such as M (2-mercaptophenothiazole), DM (dibenzothiazylsulfide) and CZ (N-cyclhexyl-2-benzothiazylsulfenamide) as well as guanidines such as DPG (diphenylguanidine), any of which may be used in an amount of 0.1 to 5 parts by weight, preferably 0.2 to 3 parts by weight per 100 parts by weight of the rubber material.

[0061] This rubber composition may contain, in addition to those described above, additives usually employed in 15 the rubber industries such as anti-aging agent, zinc oxide, stearic acid, antioxidants and anti-ozone degradation agent.

[0062] In the method according to the present invention, after incorporating the components described above, a kneader such as a roll and an internal mixer may be used to yield a rubber composition.

[0063] The rubber composition thus obtained may be molded and vulcanized and then used in tire components such 20 as tire treads, under treads, carcasses, side walls and bead parts, as well as in other industrial materials such as vibration proof rubbers, belts and hoses. In particular, it is used preferably as a rubber for a tire tread.

20 Examples

[0064] The present invention will be further described in the following examples, which are not intended to restrict the present invention.

[0065] In the examples, parts and percents are ones by weight unless otherwise specified.

[0066] Measurements were conducted as follows.

[0067] The microstructure of the butadiene moiety of a polymer was determined by infrared method (Morero method). The bound styrene content in a butadiene/styrene copolymer was calculated based on the ratio of integrated values from ¹H-NMR spectrum.

[0068] Compound Mooney (ML₁₊₄) was determined at 128°C according to JIS K6300.

[0069] As an index of hysteresis loss, tan δ was employed. A composition having a lower tan δ was regarded as that having a lower hysteresis loss. The value of tan δ was determined using a rheometer (Rheometrics) at the temperature of 50 °C with the strain of 8% and the frequency of 15 Hz.

[0070] Physical property (tensile strength at break, extension at break) and 300 % modulus (M₃₀₀) were determined 35 according to JIS K6301.

[0071] Wearing resistance was determined by measuring an amount of wearing at the slip rate of 60 % at a room temperature using a Lambourn type wearing tester. Wearing resistance index was represented as a value based on the wearing resistance of the vulcanized rubber obtained from Polymer C in Comparative Example 1 being 100. A greater index reflects a better result.

40 Example 1

Synthesis of Polymer A and preparation of Master batch A

[0072] In a 5 liter reaction vessel which was dried and purged with nitrogen, 2000 g of cyclohexane, 400 g of 1,3-butadiene monomer, 100 g of styrene monomer and 10 g of tetrahydrofuran were charged and then the solution was heated to 40 °C, and 0.32 g of n-butyllithium was added thereto to initiate polymerization. After checking the peak temperature of the polymerization and reduction in pressure, γ-glycidoxypropyltrimethoxysilane (GPMOS) as a modifier was added in the amount equimolar to n-butyllithium (1.18g), and then the reaction was further proceeded for 30 minutes.

[0073] About a half of the copolymer solution was taken as a sample, and an antioxidant di-t-butyl-p-cresol was added thereto in the amount of 0.5 g per 100 g of the copolymer. Then the solution was dried to remove the solvent in a conventional manner to obtain the copolymer composition designated as Polymer A.

[0074] To 100 parts by weight of the remainder of the copolymer solution, 50 parts by weight of a silica (VN3: Tradename, Nippon Silica) was admixed and the mixture was stirred at a room temperature for about 1 hour thereby reacting the terminals of the copolymer with the silica. Finally, di-t-butyl-p-cresol was added thereto in the amount of 0.5 g per 100 g of the copolymer, and the solution was dried to remove the solvent in a conventional manner to obtain the terminally-modified copolymer composition designated as Master batch A.

[0075] The microstructure of this copolymer was represented by the styrene content of 20 % by weight and the vinyl content of 50 % by weight for the total butadiene unit.

Example 2

5

Synthesis of Polymer B and preparation of Master batch B

[0076] Polymer B and Master batch B were obtained similarly as in Example 1 except for adding 1.24 g of γ -methacryloxypropyl trimethoxysilane (γ -MS) as a modifier instead of CPMOS.

10

[0077] The microstructure of this copolymer was represented by the styrene content of 21 % by weight and the vinyl content of 49 % by weight.

Comparative Example 1

15

Synthesis of Polymer C and preparation of Master batch C

[0078] Polymer C and Master batch C were obtained similarly as in Example 1 except for adding no modifier GPMOS.

[0079] The microstructure of this copolymer was represented by the styrene content of 21 wt.% and the vinyl content of 51 wt.%.

20

[0080] To each of the polymers and master batches obtained as described above, various additives were incorporated in the amounts shown in Table 1, and kneaded and then vulcanized for 30 minutes at 140 °C to prepare a vulcanized rubber. The physical properties of the vulcanized rubbers thus obtained were evaluated and the results are shown in Table 2.

25

Comparative Example 2

Synthesis of Polymer D and preparation of Master batch D

30

[0081] Polymer D and Master batch D were obtained similarly as in Example 1 except for adding 0.89 g of methyltriethoxysilane instead of GPMOS.

[0082] The microstructure of this copolymer was represented by the styrene content of 20 wt.% and the vinyl content of 51 wt.%.

35

[0083] To each of the polymers and master batches obtained as described above, various additives were incorporated in the amounts shown in Table 1, and kneaded and then vulcanized for 30 minutes at 140 °C to prepare a vulcanized rubber. The physical properties of the vulcanized rubbers thus obtained were evaluated and the results are shown in Table 2.

Table 1

40	Polymer + silica or Master batch	150*1 (parts by weight)
	Stearic acid	2
	Antioxidant *2	1
	Zinc oxide	3
	Aromatic oil	15
45	Vulcanization accelerator (DPG*3)	0.8
	Vulcanization accelerator (DM*4)	1.2
	Sulfur	2.0

*1 Polymer 100 parts + silica 50 parts

*2 Di-t-butyl-p-cresol

*3 Diphenylguanidine

*4 Dibenzothiazyl disulfide

55

Table 2

	Examples				Comparative Examples			
	Master Batch A	Master Batch B	Polymer A	Polymer B	Master Batch C	Polymer C	Master Batch D	Polymer D
Mooney viscosity (ML _{1,4})	65	69	72	76	88	90	73	75
Tensile strength break characteristics (kg/cm ²)	220	205	250	240	181	180	210	200
Extension (%)	490	475	450	440	530	550	500	510
300 % Modulus (kg/cm ²)	90	87	100	98	62	58	78	74
$\tan \delta$ 50 °C	0.09	0.09	0.07	0.08	0.13	0.14	0.10	0.10
Wearing resistance index	119	117	120	118	101	100	112	110

[0084] As evident from Table 2, the rubber compositions obtained by the present method had excellent break characteristics, modulus, wearing resistance and low hysteresis loss ($\tan \delta : 50^\circ \text{ C}$) as well as well-balanced physical properties including processability, while they exhibited increased values of compound Mooney.

[0085] The vulcanized rubbers obtained from Polymers A and B which were the copolymers before incorporation of the silica also exhibited excellent properties as shown in Table 2, and the vulcanized rubbers obtained from Master batches A and B after incorporation of the silica exhibited further improved properties, indicating that the disadvantages involved in the conventional silica incorporation were eliminated.

[0086] In addition, as compared with Comparative Examples master batch C and polymer C in which any modifier to be used in the present invention was not employed, master batches of the Examples described above had lower Mooney viscosity and yielded vulcanized rubbers having higher tensile strength and lower $\tan \delta$, indicating excellent processability and break characteristics and satisfactorily low hysteresis loss.

[0087] Also as compared with Comparative Examples master batch D and polymer D in which a conventional modifier was used, the Examples exhibited improvement in all items of processability, break characteristics and low hysteresis loss, and were especially excellent with regard to break characteristics.

[0088] According to the method for producing a rubber composition according to the present invention, a rubber composition can be obtained which has high wet grip characteristics, break characteristics and wearing resistance and satisfactorily low hysteresis loss (low rolling resistance, low gas mileage).

20 Claims

1. A method for producing a rubber composition which comprises:

25 the step wherein a conjugated diene monomer, or a conjugated diene monomer and a vinyl aromatic hydro-carbon monomer are polymerized by using an organic lithium initiator;
 the step wherein an active terminal of the polymer or copolymer thus obtained is modified with a modifier represented by the general formula (I) shown below and having reactivity with a silica surface; and
 the step wherein the terminally-modified polymer or terminally-modified copolymer thus obtained is mixed with a silica in an organic solvent;



35 wherein R^1 denotes a substituent having an epoxy group or an unsaturated carbonyl group, X denotes an alkyl group or a halogen atom, R^2 denotes a group selected from the group consisting of aliphatic, alicyclic and aromatic hydrocarbons of 1 to 20 carbon atoms, m denotes an integer from 1 to 3, n denotes an integer from 1 to 3 and $m+n$ is an integer from 2 to 4.

40 2. A method for producing a rubber composition according to Claim 1, wherein R^1 denotes a substituent of 3 to 20 carbon atoms having an epoxy group or an unsaturated carbonyl group.

45 3. A method for producing a rubber composition according to Claim 2, wherein R^1 denotes a group selected from the group consisting of γ -glycidoxethyl, γ -glycidoxpropyl, γ -glycidoxbutyl, γ -methacryloxyethyl, γ -methacryloxyethyl, γ -methacryloxypropyl, β -(3,4-epoxycyclohexyl)ethyl and β -(3,4-epoxycyclohexyl)propyl.

4. A method for producing a rubber composition according to any one of Claims 1-3, wherein X is an alkyl group of 1 to 20 carbon atoms or a halogen atom.

50 5. A method for producing a rubber composition according to any one of Claims 1-4, wherein R^2 is a hydrocarbon group selected from the group consisting of aliphatic, alicyclic and aromatic groups of 1 to 13 carbon atoms.

6. A method for producing a rubber composition according to Claim 1, wherein said modifier having reactivity with a silica surface is γ -glycidoxpropyl trimethoxysilane or γ -methacryloxypropyl trimethoxysilane.

55 7. A method for producing a rubber composition according to any one of Claims 1-6, wherein the amount of the organic lithium initiator used is from 0.2 to 30 mmol per 100 g of the monomer, the amount of the modifier used is from 0.1 to 10 molecules per lithium atom in the organic lithium initiator, and the amount of the silica incorporated is from 1 to 100 parts by weight per 100 parts by weight of said terminally-modified polymer or terminally modified

copolymer.

8. A method for producing a rubber composition according to any one of Claims 1-7, wherein the silica is a wet silica.
- 5 9. A method for producing a rubber composition according to any one of Claims 1-8 wherein the rubber composition has a Mooney viscosity (ML₁₊₄, 100 ° C) within the range from 20 to 200.
10. A method for producing a rubber composition which comprises :

10 the step wherein a conjugated diene monomer of 4 to 12 carbon atoms, or said conjugated diene monomer and aromatic vinyl monomer are polymerized by using an organic lithium initiator;

the step wherein a polymeric active terminal of the polymer or copolymer thus obtained is modified with a modifier represented by the general formula (Ia) shown below and having reactivity with a silica surface; and,

15 the step wherein the terminally-modified polymer or terminally-modified copolymer thus obtained is mixed with a silica in an organic solvent;



20 wherein R¹ denotes a group selected from the group consisting of γ -glycidoxethyl, γ -glycidoxpropyl, γ -glycidoxbutyl, γ -methacryloxymethyl, γ -methacryloxyethyl, γ -methacryloxypropyl, β -(3,4-epoxycyclohexyl)ethyl and β -(3,4-epoxycyclohexyl)propyl, X¹ denotes an alkyl group of 1 to 20 carbon atoms or a halogen atom, R² denotes a group selected from the group consisting of methyl, ethyl, propyl, butyl, phenyl, naphthyl, biphenyl, anthryl and phenanthryl, m denotes an integer from 1 to 3, n denotes an integer from 1 to 3 and m+n is an integer from 2 to 4.

Patentansprüche

- 30 1. Verfahren zur Herstellung einer Kautschukzusammensetzung, umfassend
 - den Schritt, in dem ein konjugiertes Dien-Monomer oder ein konjugiertes Dien-Monomer und ein Vinyl- aromatisches Kohlenwasserstoff-Monomer unter Verwendung eines organischen Lithium-Initiators polymerisiert wird (werden);
 - den Schritt, in dem ein aktives Ende des so erhaltenen Polymers oder Copolymers mit einem Modifikations- mittel, das durch die unten angegebene allgemeine Formel (I) dargestellt wird und das hohe Reaktivität zu einer Siliciumdioxid-Oberfläche hat, modifiziert wird; und
 - den Schritt, in dem das so erhaltene am Ende modifizierte Polymer oder am Ende modifizierte Copolymer mit einem Siliciumdioxid in einem organischen Lösungsmittel vermischt wird:



45 worin R¹ für einen Substituenten, der eine Epoxy-Gruppe oder eine ungesättigte Carbonyl-Gruppe hat, steht, X eine Alkyl-Gruppe oder ein Halogenatom darstellt, R² für eine Gruppe, die aus der Gruppe, bestehend aus aliphatischen, alicyclischen und aromatischen Kohlenwasserstoffen mit 1 bis 20 Kohlenstoffatomen, ausgewählt wird, steht, m eine ganze Zahl von 1 bis 3 bedeutet, n eine ganze Zahl von 1 bis 3 bedeutet und m+n eine ganze Zahl von 2 bis 4 ist.

- 50 2. Verfahren zur Herstellung einer Kautschukzusammensetzung nach Anspruch 1, wobei R¹ für einen Substituenten mit 3 bis 20 Kohlenstoffatomen, der eine Epoxy-Gruppe oder eine ungesättigte Carbonyl-Gruppe hat, steht.
- 55 3. Verfahren zur Herstellung einer Kautschukzusammensetzung nach Anspruch 2, wobei R¹ für eine Gruppe steht, die aus der Gruppe, bestehend aus γ -Glycidoxethyl, γ -Glycidoxpropyl, γ -Glycidoxbutyl, γ -Methacryloxymethyl, γ -Methacryloxyethyl, γ -Methacryloxypropyl, β -(3,4-Epoxy- cyclohexyl)ethyl und β -(3,4-Epoxy- cyclohexyl)propyl, ausgewählt wird.

4. Verfahren zur Herstellung einer Kautschukzusammensetzung nach einem der Ansprüche 1 bis 3, wobei X eine Alkyl-Gruppe mit 1 bis 20 Kohlenstoffatomen oder ein Halogenatom ist.

5. Verfahren zur Herstellung einer Kautschukzusammensetzung nach einem der Ansprüche 1 bis 4, wobei R² eine Kohlenwasserstoff-Gruppe ist, die aus der Gruppe, bestehend aus aliphatischen, alicyclischen und aromatischen Gruppen mit 1 bis 13 Kohlenstoffatomen, ausgewählt wird.

10. Verfahren zur Herstellung einer Kautschukzusammensetzung nach Anspruch 1, wobei das Modifikationsmittel, das Reaktivität zu einer Siliciumoxid-Oberfläche hat, γ -Glycidoxypyropyltrimethoxysilan oder γ -Methacryloxypropyltrimethoxysilan ist.

15. Verfahren zur Herstellung einer Kautschukzusammensetzung nach einem der Ansprüche 1 bis 6, wobei die Menge des organischen Lithium-Initiators, die verwendet wird, 0,2 bis 30 mmol pro 100 g Monomer ist, die Menge des Modifikationsmittels, die verwendet wird, 0,1 bis 10 Moleküle pro Lithiumatom in dem organischen Lithium-Initiator ist und die Menge an eingearbeitetem Siliciumdioxid 1 bis 100 Gew.-Teile pro 100 Gew.-Teile des am Ende modifizierten Polymers oder am Ende modifizierten Copolymers ist.

20. Verfahren zur Herstellung einer Kautschukzusammensetzung nach einem der Ansprüche 1 bis 7, wobei das Siliciumdioxid ein nasses Siliciumdioxid ist.

25. Verfahren zur Herstellung einer Kautschukzusammensetzung nach einem der Ansprüche 1 bis 8, wobei die Kautschukzusammensetzung eine Mooney-Viskosität (ML₁₊₄, 100°C) im Bereich von 20 bis 200 hat.

10. Verfahren zur Herstellung einer Kautschukzusammensetzung, umfassend

25. den Schritt, in dem ein konjugiertes Dien-Monomer mit 4 bis 12 Kohlenstoffatomen oder dieses konjugierte Dien-Monomer und aromatisches Vinyl-Monomer unter Verwendung eines organischen Lithium-Initiators polymerisiert wird (werden);

30. den Schritt, in dem ein aktives Polymerende des so erhaltenen Polymers oder Copolymers mit einem Modifikationsmittel, das durch die unten angegebene allgemeine Formel (Ia) dargestellt wird und das hohe Reaktivität zu einer Siliciumdioxid-Oberfläche hat, modifiziert wird; und

35. den Schritt, in dem das so erhaltene am Ende modifizierte Polymer oder am Ende modifizierte Copolymer mit einem Siliciumdioxid in einem organischen Lösungsmittel vermischt wird:



40. worin R'¹ für eine Gruppe steht, die aus der Gruppe, bestehend aus γ -Glycidoxypyropyl, γ -Glycidoxypyropyl, γ -Glycidoxypyropyl, γ -Methacryloxyethyl, γ -Methacryloxyethyl, γ -Methacryloxyethyl, β -(3,4-Epoxyhexyl)ethyl und β -(3,4-Epoxyhexyl)ethyl, ausgewählt wird, X' eine Alkyl-Gruppe mit 1 bis 20 Kohlenstoffatomen oder ein Halogenatom darstellt, R'² für eine Gruppe, die aus der Gruppe bestehend aus Methyl, Ethyl, Propyl, Butyl, Phenyl, Naphthyl, Biphenyl, Anthryl und Phenanthryl, ausgewählt wird, steht, m eine ganze Zahl von 1 bis 3 bedeutet, n eine ganze Zahl von 1 bis 3 bedeutet und m+n eine ganze Zahl von 2 bis 4 ist.

Revendications

50. 1. Procédé de préparation d'une composition de caoutchouc qui comprend :

une étape dans laquelle un monomère de diène conjugué, ou un monomère de diène conjugué et un monomère d'hydrocarbure aromatique vinylique sont polymérisés en utilisant un amorceur organique à base de lithium ;

55. l'étape dans laquelle une terminaison active du polymère ou du copolymère ainsi obtenu est modifiée avec un agent de modification représenté par la formule générale (I) indiquée ci-dessous et possédant un caractère réactif vis-à-vis d'une surface de silice ; et

une étape dans laquelle le polymère à terminaison modifiée ou le copolymère à terminaison modifiée ainsi

obtenu est mélangé avec une silice dans un solvant organique ;



5

dans laquelle R^1 représente un substituant contenant un groupe époxy ou un groupe carbonyle insaturé, X représente un groupe alkyle ou un atome d'halogène, R^2 représente un groupe choisi dans le groupe composé des hydrocarbures aliphatiques, alicycliques et aromatiques ayant de 1 à 20 atomes de carbone, m représente un nombre entier de 1 à 3, n représente un nombre entier allant de 1 à 3 et $m+n$ est un nombre entier allant de 2 à 4.

10

2. Procédé de préparation d'une composition de caoutchouc selon la revendication 1, dans lequel R^1 représente un substituant possédant de 3 à 20 atomes de carbone contenant un groupe époxy ou un groupe carbonyle insaturé.

15

3. Procédé de préparation d'une composition de caoutchouc selon la revendication 2, dans lequel R^1 représente un groupe choisi dans le groupe composé du γ -glycidoxyméthyle, du γ -glycidoxypropyle, du γ -glycidoxylbutyle, du γ -méthacryloxyméthyle, du γ -méthacryloxyéthyle, du γ -méthacryloxypropyle, du β -(3,4-époxypropyle)éthyle et du β -(3,4-époxypropyle)cyclohexylpropyle.

20

4. Procédé de préparation d'une composition de caoutchouc selon l'une quelconque des revendications 1 à 3, dans lequel X est un groupe alkyle ayant de 1 à 20 atomes de carbone ou un atome d'halogène.

25

5. Procédé de préparation d'une composition de caoutchouc selon l'une quelconque des revendications 1 à 4, dans lequel R^2 est un groupe hydrocarbure choisi dans le groupe composé des groupes aliphatiques, alicycliques et aromatiques possédant de 1 à 13 atomes de carbone.

30

6. Procédé de préparation d'une composition de caoutchouc selon la revendication 1, dans lequel ledit agent de modification possédant un caractère réactif vis-à-vis d'une surface de silice est un γ -glycidoxypropyltriméthoxysilane ou un γ -méthacryloxypropyltriméthoxysilane.

7. Procédé de préparation d'une composition de caoutchouc selon l'une quelconque des revendications 1 à 6, dans lequel la quantité d'amorceur organique à base de lithium utilisée est de 0,2 à 30 mmoles pour 100 g de monomère, la quantité de l'agent de modification utilisée est de 0,1 à 10 molécules par atome de lithium dans l'amorceur organique à base de lithium, et la quantité de silice incorporée est de 1 à 100 parties en poids pour 100 parties en poids dudit polymère à terminaison modifiée ou copolymère à terminaison modifiée.

35

8. Procédé de préparation d'une composition de caoutchouc selon l'une quelconque des revendications 1 à 7, dans lequel la silice est une silice hydratée.

40

9. Procédé de préparation d'une composition de caoutchouc selon l'une quelconque des revendications 1 à 8, dans lequel la composition de caoutchouc possède une viscosité Mooney (ML_{1+4} , 100°C) comprise dans la plage allant de 20 à 200.

45

10. Procédé de préparation d'une composition de caoutchouc qui comprend :

une étape dans laquelle un monomère de diène conjugué possédant de 4 à 12 atomes de carbone, ou ledit monomère de diène conjugué et un monomère vinylique aromatique sont polymérisés en utilisant un amorceur organique à base de lithium ;

50

une étape dans laquelle une terminaison polymère active du polymère ou du copolymère ainsi obtenu est modifiée avec un agent de modification représenté par la formule générale (la) indiquée ci-dessous et possédant un caractère réactif vis-à-vis une surface de silice ; et

une étape dans laquelle le polymère à terminaison modifiée ou le copolymère à terminaison modifiée ainsi obtenu est mélangé avec une silice dans un solvant organique ;

55



dans laquelle R^1 représente un groupe choisi dans le groupe composé du γ -glycidoxyméthyle, du γ -gly-

cidoxypropyle, du γ -glycidoxypbutyle, du γ -méthacryloxyméthyle, du γ -méthacryloxypropyle, du β - (3,4-époxy-cyclohexyl)éthyle et du β -(3,4-époxy-cyclohexyl)propyle, X' représente un groupe alkyle possédant de 1 à 20 atomes de carbone ou un atome d'halogène, R² représente un groupe choisi dans le groupe composé des groupes méthyle, éthyle, propyle, butyle, phényle, naphtyle, biphenyle, anthryle et phénanthryle, m représente un nombre entier allant de 1 à 3, n représente un nombre entier de 1 à 3 et m+n est un nombre entier allant de 2 à 4.

5

10

15

20

25

30

35

40

45

50

55